PATENT Docket No. SALK1650-2 (088802-2753)

## THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in this application.

- 1. (Previously presented) A method for treating an individual suffering from diabetes mellitus, said method comprising administering an effective amount of a compound which inhibits binding of CREB to CBP.
- 2 (Original) A method according to claim 1 wherein said treatment of diabetes mellitus ameliorates hyperglycemia.
- 3. (Original) A method according to claim 2 wherein gluconeogenesis is modulated.
- 4. (Original) A method according to claim 3 wherein transcription of PEPCK is //inhibited.
- 5. (Previously presented) A method according to claim 2 wherein transcription of the glucagon gene is inhibited.
- 6. (Previously presented) A method according to claim 1 wherein said individual is a human.
- 7. (Previously presented) A method according to claim 1 wherein said administering is accomplished by oral, intravenous, subcutaneous, intramuscular or intracutaneous mode of administration.
- 12. (Previously presented) A method for treating an individual suffering from diabetes mellitus, comprising administering an effective amount of a compound which disrupts complex

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comprising cyclic AMP response element binding protein (CREB) and CREB binding protein (CBP), said compound identified by a method comprising:

- (a) contacting a modified host cell with a test compound, wherein said modified host cell comprises:
  - a first fusion protein comprising a GAL4 DNA binding domain, operatively associated with the kinase-inducible domain (KID) of CREB,
  - a second fusion protein comprising an activation domain, operatively associated with the CREB binding domain (KIX) of CBP, and
  - a reporter construct comprising a GAL4 response element operatively linked to a reporter gene; and
- (b) selecting those test compounds which cause reduced expression of the reporter gene product, wherein said compounds are identified as disrupting complex comprising CREB and CBP.
- 17. (Previously presented) A method for treating an individual suffering from diabetes mellitus, comprising administering an effective amount of a compound which disrupts complex comprising cyclic AMP response element binding protein (CREB) and CREB binding protein (CBP), said compound identified by a method comprising:
- (a) contacting a modified host cell with a test compound, wherein said modified host cell comprises:
  - a first fusion protein comprising an activation domain, operatively associated with the kinase-inducible domain (KID) of CREB,
  - a second fusion protein comprising a GAL4 DNA binding domain operatively associated with the CREB binding domain (KIX) of CBP, and
  - a reporter construct comprising a GAL4 response element operatively linked to a reporter gene; and



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- (b) selecting those test compounds which cause reduced expression of the reporter gene product, wherein said compounds are identified as disrupting complex comprising CREB and CBP.
- 18. (Previously presented) A method for modulating glucose metabolism in an individual, said method comprising administering an effective amount of a compound which inhibits binding of CREB to CBP.
- 19. (Previously presented) A method according to claim 18 wherein said modulating glucose metabolism results in decreased serum glucose.
- 20. (Previously presented) A method according to claim 18 wherein said modulating glucose metabolism results in decreased gluconeogenesis.
- 21. (Previously presented) A method according to claim 20 wherein transcription of PEPCK is inhibited.
- 22. (Previously presented) A method according to claim 20 wherein transcription of the glucagon gene is inhibited.
- 23. (Previously presented) A method according to claim 18 wherein said individual is a human.
- 24. (Previously presented) A method according to claim 18 wherein said administering is accomplished by oral, intravenous, subcutaneous, intramuscular or intracutaneous mode of administration.

- 25. (Previously presented) A method for inhibiting expression of phosphoenolpyruvate carboxykinase (PEPCK) enzyme in an individual, said method comprising administering an effective amount of a compound which inhibits binding of CREB to CBP.
- 26. (Previously presented) A method according to claim 25 wherein said inhibiting PEPCK enzyme expression results in decreased serum glucose.
- 27. (Previously presented) A method according to claim 25 wherein said inhibiting PEPCK enzyme expression results in decreased gluconeogenesis.
- 28. (Previously presented) A method according to claim 27 wherein transcription of PEPCK is inhibited.
- 29. (Previously presented) A method according to claim 27 wherein transcription of the glucagon gene is inhibited.
- 30. (Previously presented) A method according to claim 25 wherein said individual is a human.
- 31. (Previously presented) A method according to claim 30 wherein said individual is suffering from diabetes mellitus.
- 32. (Previously presented) A method according to claim 25 wherein said administering is accomplished by oral, intravenous, subcutaneous, intramuscular or intracutaneous mode of administration.
- 33. (Previously presented) A method according to claim 23 wherein said individual is suffering from diabetes mellitus.